

**TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371**

20-4736P
U.S. APPLICATION NO. (if known, see 37 CFR 1.5)
09762284

INTERNATIONAL APPLICATION NO.

PCT/JP99/00638

INTERNATIONAL FILING DATE

February 15, 1999

PRIORITY DATE CLAIMED

February 15, 1998

TITLE OF INVENTION

NOVEL PROPHYLACTICS/REMEDIES FOR IMMUNOPATHY

APPLICANT(S) FOR DO/EO/US

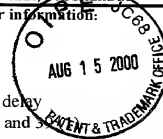
SASAKI, Takumi; KIMACHI, Kazuhiko; SOEJIMA, Kenji; NOZAKI, Chikateru; KIMURA, Yumi; FUJIYAMA, Yoshitaka

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 33.
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(3)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(2)).
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.-1449 and International Search Report (PCT/ISA/210)
12. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A FIRST preliminary amendment.
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:
 - 1.) Seven (7) sheets of Formal Drawings
 - 2.) International Preliminary Examination Report



U.S. APPLICATION NO. 097622284 NEW	INTERNATIONAL APPLICATION NO. PCT/JP99/00638	534 Rec'd PCT/PTO 15 AUG 2000 ATTORNEY'S DOCKET NUMBER 20-4736P												
17. <input checked="" type="checkbox"/> The following fees are submitted: BASIC NATIONAL FEE (37 CFR 1.492(a)(1)-(5): Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO. \$970.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$840.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO. \$690.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$670.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4). \$96.00 ENTER APPROPRIATE BASIC FEE AMOUNT =		<table border="1" style="width:100%; border-collapse: collapse;"> <tr> <th colspan="2" style="text-align: left;">CALCULATIONS</th> <th style="text-align: left;">PTO USE ONLY</th> </tr> <tr> <td style="width:50%;"></td> <td style="width:25%;"></td> <td style="width:25%;"></td> </tr> <tr> <td>\$ 840.00</td> <td></td> <td></td> </tr> <tr> <td>\$ 0</td> <td></td> <td></td> </tr> </table>	CALCULATIONS		PTO USE ONLY				\$ 840.00			\$ 0		
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Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).														
CLAIMS	NUMBER FILED	NUMBER EXTRA												
Total Claims	24 - 20 =	4												
Independent Claims	1 - 3 =	0												
MULTIPLE DEPENDENT CLAIM(S) (if applicable) Yes		+ \$260.00												
TOTAL OF ABOVE CALCULATIONS =		\$ 1172.00												
Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28).		\$ 0												
SUBTOTAL =		\$ 1172.00												
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).		\$ 0												
TOTAL NATIONAL FEE =		\$ 1172.00												
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +		\$ 40.00												
TOTAL FEES ENCLOSED =		\$ 1212.00												
		Amount to be: refunded \$												
		charged \$												

a. ☒ A check in the amount of \$ 1212.00 to cover the above fees is enclosed.

b. ☐ Please charge my Deposit Account. No. _____ in the amount of \$ _____ to cover the above fees.
 A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any
 overpayment to Deposit Account No. 02-2448.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR
 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

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 #32.181 (MSW)
 REGISTRATION NUMBER

09/622284

534 Rec'd PCT/PTO 15 AUG 2000

PATENT
20-4736P

IN THE U.S. PATENT AND TRADEMARK OFFICE

Applicant: SASAKI, Takumi et al.
Int'l. Appl. No.: PCT/JP99/00638
Appl. No.: New Group:
Filed: August 15, 2000 Examiner:
For: NOVEL PROPHYLACTICS/REMEDIES FOR
IMMUNOPATHY

PRELIMINARY AMENDMENT

BOX PATENT APPLICATION

Assistant Commissioner for Patents
Washington, DC 20231

August 15, 2000

Sir:

The following Preliminary Amendments and Remarks are respectfully submitted in connection with the above-identified application.

AMENDMENTS

IN THE SPECIFICATION:

Please amend the specification as follows:

Before line 1, insert --This application is the national phase under 35 U.S.C. § 371 of PCT International Application No. PCT/JP99/00638 which has an International filing date of February 15, 1999, which designated the United States of America.--

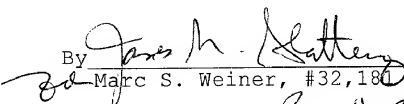
REMARKS

The specification has been amended to provide a cross-reference to the previously filed International Application.

If necessary, the Commissioner is hereby authorized in this, concurrent, and future replies, to charge payment or credit any overpayment to Deposit Account No. 02-2448 for any additional fees required under 37 C.F.R. § 1.16 or under 37 C.F.R. § 1.17; particularly, extension of time fees.

Respectfully submitted,

BIRCH, STEWART, KOLASCH & BIRCH, LLP

By 

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MSW/cqc
20-4736P

(Rev. 04/19/2000)

DESCRIPTION

NOVEL PROPHYLACTICS/REMEDIES FOR IMMUNOPATHY

TECHNICAL FIELD OF THE INVENTION

5 The present invention relates to novel
prophylactics/remedies for immunopathy. More specifically,
the present invention relates to novel
prophylactics/remedies for immunopathy such as rheumatoid
arthritis, allergic diseases, etc. comprising as an active
ingredient modifications of natural Staphylococcal
10 enterotoxin B (hereinafter also referred to as "SEB"), known
as one of superantigens, or derivatives thereof.

BACKGROUND OF THE INVENTION

Autoimmune diseases are classified into two types:
organ-nonspecific type autoimmune diseases such as
15 rheumatoid arthritis (hereinafter also referred to as "RA")
and organ-specific type autoimmune diseases such as
ulcerative colitis. They are induced by T cells responsive
to self antigens, said T cells being normally under
immunological tolerance, that were activated within self
20 tissues by some reasons to respond to self antigens, leading
to continuous inflammatory reactions to thereby damage
tissues. In such cases, self antigens are type II collagen
that constitutes self joint or main components of the mucous
membrane of the intestine, respectively.

25 The number of patients suffering from these

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diseases has slightly increased year by year but no effective remedies or prophylaxis have been found ("Immunodeficiency due to medicament", Men-eki Kagaku (Immunological Science), Vol. 9. p.285-289 (1984) Ed. by Yuichi Yamamura, Chuzo Kishimoto, Robert A. Good).

Currently, for treatment of these diseases, there have been employed pharmacotherapy including administration of Salazopyrin, 5-aminosalicylic acid, azathioprine, 6-MP, tranilast, methotrexate, cyclosporine A, or metronidazole, and administration of an excess amount of 7S-immunoglobulin; surgical therapy such as thymectomy or replacement with artificial joint; or symptomatic therapy such as nutritional therapy (Yoichi Ichikawa et al. "Study on efficacy of long-term administration of methotrexate and salazosulfapyridine on rheumatoid arthritis case" Rheumatism, Vol. 35, p.663-670, (1995); Sadao Kashiwazaki, "Study on efficacy of combination of auranofin and methotrexate on rheumatoid arthritis", Rheumatism, Vol. 36, p.528-544, (1996); Takefumi Furutani et al., "Detrimental event in therapy with low dose methotrexate on rheumatoid arthritis", Rheumatism, Vol. 36, p.746-752, (1996); Nobuo Watanabe, "Pharmacotherapy on juvenile rheumatoid arthritis", Rheumatism, Vol. 36, p.670-675, (1996); Takayasu Yakura, "Immunosuppressive therapy: Treatment of autoimmune diseases", Sogo Rinsho, Vol. 30, p.3358, (1981); Shin Totokawa et al., "Study on methotrexate

therapy in rheumatoid arthritis: Seeking for strategy of more effective administration", Rheumatism, Vol. 37, p.681-687 (1997)). However, these therapies are not eradicated but rather are disadvantageous in that they may cause severe adverse side effects due to long-term ingestion of medicaments. Thus, it is desired to develop more effective prophylactics/remedies and therapy.

As is well known, SEB is one of bacterial superantigens (White J. et al., Cell, Vol. 56, p.27-35, 1989). Normal antigens, being complexed with class II Major Histocompatibility Complex (hereinafter also referred to as "MHC"), are recognized by T cell antigen receptor (hereinafter also referred to as "TCR"). This recognition is restricted to a haplotype of class II MHC, called "MHC restriction". On the contrary, superantigens are bound to class II MHC molecule irrespective of haplotype and further to specific β chain variable region (V β chain) of TCR. When such a linkage is formed, T cells bound with the superantigen are transiently activated, are promoted to divide and propagate and produce inflammatory cytokines (Micusan V.V. & Thibodean J., Seminars in Immunology, Vol. 5, p.3-11, 1993).

SEB is one of toxins causing food poisoning. Symptoms of food poisoning can be seen when SEB is intravenously administered to animals, suggesting that

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toxicity is exerted through the above-mentioned biological activity of SEB (Micusan V.V. & Thibodean J., Seminars in Immunology, Vol. 5, p.3-11, 1993).

However, when a superantigen is intravenously or
5 intraperitoneally administered to newborn mice, a subpopulation of T cells having a V β chain responsive to the superantigen is eliminated so that said mice become non-responsive to said antigen, i.e. immunological tolerance (White J. et al., Cell, Vol. 56, p.27-35, 1989). On the
10 other hand, when SEB is administered to adult mice, after transient activation as mentioned above, a subpopulation of T cells having responsive V β 7, V β 8 TCR becomes non-responsive to another stimulation with SEB, i.e. becomes anergy, to thereby cause immunological tolerance. With such
15 activity, SEB is suggested to be potential prophylactics/remedies for intractable autoimmune diseases such as rheumatoid arthritis or ulcerative colitis or immunopathy (Japanese Patent Publication No. 9-110704). Since SEB is an enterotoxin causing staphylococcal food
20 poisoning, pathogenicity of SEB is a problem when it is administered into the living body. In accordance with Japanese Patent Publication No. 9-110704, such toxicity is avoided by continuously administering a highly purified SEB in such a dose that does not render SEB being pathogenic
25 through oral route for a long period of time to thereby

effectively induce immunological tolerance without pathogenicity of SEB.

Another approach for avoiding SEB toxicity is "protective action of mutated superantigen" as disclosed by Kappler J. and Marrack P. et al. in Japanese Patent Publication No. 8-500328. Their invention is summarized as follows: SEB mutants are prepared and purified and those SEB mutants that could bind to both human class II MHC molecule and TCR were selected. Then, the SEB mutants, that bound to class II-positive cells to a degree undistinguishable from that of natural type SEB but did not stimulate propagation of T cells, were injected to an animal prior to exposition to SEB. As a result, the injected animal exhibited complete protection from toxicity of SEB.

The most important teaching of Japanese Patent Publication No. 8-500328 is a study on a structure of SEB that can be utilized for elimination of toxicity. A three-dimensional structure of SEB was reported by Swaminathan S. et al. (Nature, Vol. 359, p.801 (1992)). SEB molecule is consisted of two domains, i.e. the first domain consisted of residues 1 to 120 and the second consisted of residues 127-239. At the N-terminal of SEB, there have been identified three regions, i.e. Region 1 (residues 9 to 23), Region 2 (residues 41 to 53) and Region 3 (residues 60 to 61), that may affect class II MHC cell binding and/or TCR binding.

DISCLOSURE OF THE INVENTION

Based on the above findings, the present inventors have completed the present invention that provides novel prophylactics/remedies for immunopathy comprising as an active ingredient SEB modifications that can effectively be used for prophylaxis and treatment of immunopathy with reduced toxicity or derivatives thereof.

That is, the present invention provides prophylactics/remedies for immunopathy comprising, as an active ingredient, modifications of Staphylococcal enterotoxin B (SEB) with substitution of at least one amino acid residues within the amino acid sequence of natural type SEB, or derivatives thereof, wherein said SEB modifications or derivatives thereof have inhibitory activity on T cell activation wherein they interact with specific V β component of T cell receptor (TCR) but are reduced in their immunological responsiveness to SEB without inducing elimination of T cells having specific V β component, the elimination being normally induced by natural type SEB or recombinant wild-type SEB.

In the first aspect of the present invention, the SEB modifications or derivatives thereof are SEB having amino acid substitution at the 9-position on the basis of natural type SEB with a protein-constituting amino acid other than aspartic acid, preferably asparagine, through

site-specific mutagenesis, or derivatives thereof.

In the second aspect of the present invention, the SEB modifications or derivatives thereof are SEB having amino acid substitution at the 23-position on the basis of natural type SEB with a protein-constituting amino acid other than asparagine, preferably tyrosine, aspartic acid, isoleucine or lysine, through site-specific mutagenesis, or derivatives thereof.

In the third aspect of the present invention, the SEB modifications or derivatives thereof are SEB having amino acid substitution at the 41-position on the basis of natural type SEB with a protein-constituting amino acid other than isoleucine, preferably arginine or threonine, plus amino acid substitution either at the 44-position with a protein-constituting amino acid other than phenylalanine, preferably valine, or at the 45-position with a protein-constituting amino acid other than leucine, preferably valine, through site-specific mutagenesis, or derivatives thereof.

In the fourth aspect of the present invention, the SEB modifications or derivatives thereof are SEB having amino acid substitution at the 44-position on the basis of natural type SEB with a protein-constituting amino acid other than phenylalanine, preferably serine, through site-specific mutagenesis, or derivatives thereof.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 shows nucleotide sequences of each of PCR primers used for amino acid substitution for preparing each of SEB modifications of the present invention.

5 Fig. 2 shows nucleotide sequences of each of PCR primers used for amino acid substitution for preparing each of SEB modifications of the present invention.

10 Fig. 3 shows an amount of TNF- α produced by human peripheral blood mononuclear cells (PBMC) stimulated with wild-type SEB and SEB modifications.

Fig. 4 shows T cell inhibitory activity of SEB modifications.

15 Fig. 5 shows therapeutic effects of oral administration of SEB modifications on collagen-induced arthritis for effective cases wherein the axis of ordinate indicates severity of test mice. The axis of ordinates indicates severity of the tested mice and the axis of abscissas indicates days after initiation of administration of SEB modifications.

20 Fig. 6 shows therapeutic effects of oral administration of SEB modifications on collagen-induced arthritis for non-effective cases. The axis of ordinates indicates severity of the tested mice and the axis of abscissas indicates days after initiation of administration of SEB modifications.

25

Fig. 7 shows prophylactic effects of SEB modifications orally administered on collagen-induced arthritis. The axis of ordinate indicates severity of the tested mice and the axis of abscissas indicates days after initiation of administration of SEB modifications.

BEST MODE FOR CARRYING OUT THE INVENTION

In accordance with the present invention, for preparing SEB modifications, those modifications that affect binding with class II MHC molecule were thoroughly investigated while there was introduced no modification at the sites involved in binding with TCR. Since linkage between superantigens and class II MHC molecule is indispensable to T cell activation by superantigens, it was expected that inhibition of SEB binding to class II MHC molecule would reduce responsiveness of T cells to SEB, i.e. anergy is induced without modification on TCR binding sites.

For Region 1 defined as amino acid residues Nos. 9 to 23, modifications of asparagine at the 23-position that block binding with class II MHC molecule were particularly investigated, including N23D, N23K, N23Y, and N23I. Modifications of aspartic acid at the 9-position such as D9N and of phenylalanine at the 17-position such as F17S were also investigated. The indication used herein for modifications is such that amino acids before and after modification are indicated before and after the number of

position of amino acid residues where modifications occurred, respectively. For example, "N23D" means that "N" (Asn: asparagine) at the 23-position counted from the N-terminal is replaced with "D" (Asp: aspartic acid).

5 Region 2 defined as amino acid residue Nos. 40 to 53 is most important for mediating binding with class II MHC molecule in all the Staphylococcal enterotoxins. For Region 2, mutations at the 41-, 44-, 45-, and 53-positions were investigated. These amino acid residues are specific for
10 binding to class II MHC molecule. Modifications were prepared by introducing point mutations through site-specific mutagenesis at the residues at the 41-, 44-, 45-, and 53-positions and in addition at the 59-position. Prepared modifications were: I41T L45V, I41R F44V, F44S, F44L I53N, and F44L I53N G59W.
15

The SEB modifications as mentioned above were estimated for their safety and effectiveness. As a result, it was found that the following embodiments of SEB modifications could be used as novel prophylactics/remedies
20 for immunopathy such as rheumatoid arthritis with safe and excellent activities:

(1) SEB modifications having amino acid substitution at the 9-position on the basis of natural type SEB with a protein-constituting amino acid other than aspartic acid,
25 particularly asparagine, or derivatives thereof;

(2) SEB modifications having amino acid substitution at the 23-position on the basis of natural type SEB with a protein-constituting amino acid other than asparagine, particularly tyrosine, aspartic acid, isoleucine or lysine, or derivatives thereof;

(3) SEB modifications having amino acid substitution at the 41-position on the basis of natural type SEB with a protein-constituting amino acid other than isoleucine, particularly arginine or threonine, plus amino acid substitution either at the 44-position with a protein-constituting amino acid other than phenylalanine, particularly valine, or at the 45-position with a protein-constituting amino acid other than leucine, particularly valine, or derivatives thereof; and

(4) SEB modifications having amino acid substitution at the 44-position on the basis of natural type SEB with a protein-constituting amino acid other than phenylalanine, particularly serine, or derivatives thereof.

It was also found that these SEB modifications could exert surprisingly excellent effects when they were administered through oral route.

First, reduced toxicity of the SEB modifications was investigated with induction of weight loss being as index. Weight loss was evaluated with BALB/c mice. A PBS (phosphate buffered saline) solution of 300 µg SEB, wherein

endotoxin was removed up to a concentration that did not affect the experiment, was intraperitoneally administered to mice. Next day, weighing each mouse was started. As a result, wild type SEB and D9N showed weight loss by 7 to 9% until three days after administration whereas all the other SEB modifications did not show any weight loss.

Furthermore, the present inventors investigated these SEB modifications for their lethal toxicity induced by D-galactosamine. It was reported that administration of D-galactosamine induced drastic increase in sensitivity to SEB in mice, for example, LD50 of SEB was reduced to 2 µg/mice when D-galactosamine was administered to female BALB/c mice and 100% death was observed with administration of 20 µg/mice SEB (Miethke T. et al., J. Exp. Med., vol. 175, p.91-98 (1992)).

Using this experiment system, the present inventors investigated the SEB modifications for their lethal toxicity. As a result, wild type SEB and D9N showed lethality after 48 hours of 9/10 and 8/10, respectively, whereas I41T L45V showed 3/10, I41R F44V 1/10, F44S 0/10, F44L I53N 0/10, F44L I53N 1/10, and F44L I53N G59W 0/10, respectively. Thus, it was found that these SEB modifications had extremely reduced lethal toxicity. This expressly shows that the modifications according to the present invention not only can reduce toxicity as disclosed

in Japanese Patent Publication No. 8-500328 but also eliminate lethal toxicity.

As described above, SEB is expected to be a potential prophylactic/remedy for immunopathy. However, it is also reported that SEB per se is involved in onset of immunopathy (Omata S. et al., Cellular Immunol., Vol. 179, p.138-145 (1997)). Thus, relationship between SEB per se as well as SEB modifications of the present invention and onset of immunopathy was investigated using collagen-induced arthritis (hereinafter also referred to as "CIA"), a model of human rheumatoid arthritis. Natural type SEB and SEB modifications provided by the present invention were administered to mice with arthritis induced by administration of collagen. In case of mice administered with natural type SEB, onset rate of arthritis was considerably high and severity of the disease was also significantly high, suggesting relationship between SEB and onset of the disease. On the contrary, onset of severe arthritis was not observed when SEB modifications were administered.

Some toxicity of SEB appears to be due to a monokine, especially tumor necrosis factor- α (TNF- α), produced by SEB-stimulated leukocytes. In fact, human peripheral blood lymphocytes produce a large amount of TNF- α when reacted with SEB. However, TNF- α level produced by SEB

modifications was extremely low with stimulating activity of N23Y being the lowest.

As described above, SEB modifications exhibited lowered stimulating activity for propagation, cytokine production and the like. It is estimated that this lowered activity for stimulation is closely related to reduction of lethal toxicity in mice or loss of ability to induce arthritis. Interestingly, however, SEB modifications showed inhibitory activity to T lymphocyte activation although they had extremely reduced stimulating activity to T lymphocytes. When T cells are stimulated with antigens in the presence of antigen-presenting cells, it results in propagation of T cells or production by T cells of cytokines such as gamma-interferon (γ -IFN). However, addition of SEB modifications to this system inhibited these reactions. That is, it was found that SEB modifications not only exhibited reduced toxicity as compared to natural type SEB but also acquired additional activity to inhibit T cell activation, not shown in natural type SEB.

This inhibitory effect to T cell activation was also observed in vivo. In the above experimental system, when SEB modifications of the present invention were administered prior to administration of natural type SEB deeply involved in onset of arthritis, onset of severe arthritis could be protected, supporting excellent effects

of administration of SEB modifications.

Whether SEB modifications of the present invention could be used as remedies for autoimmune disease through oral administration was investigated using collagen-induced arthritis (CIA) system. As a result, it was found that wild type and I41T L45V, I41R F44V, F44L I53N G59W, F44S, N23Y, N23I and N23D modifications inhibited exacerbation of arthritis to confirm effectiveness of these SEB modifications. Among these modifications, N23Y was the most potent inhibitor of CIA.

Accordingly, medicaments comprising SEB modifications in accordance with the present invention can be utilized as prophylactics/remedies for rheumatoid arthritis, particularly for oral administration, with lowered toxicity and more potent effectiveness based on inhibitory activity to T cell activation as compared to conventional medicaments. It is also expected that SEB modifications of the present invention exhibit prophylactic effects to inflammatory bowel diseases as in natural type SEB as disclosed in Japanese Patent Publication No. 9-110704.

The prophylactic/remedy for immunopathy for oral administration of the present invention is characterized by that it contains an extremely lowered amount of SEB modifications as compared to medicaments for another route of administration. These SEB modifications, like SEB,

specifically induce weak tolerance to a subpopulation of T cells with SEB-binding V β TCR. On the other hand, they inhibit inflammatory activity of T cells when administered orally.

5 For preparing SEB modifications for use in the present invention, any method may be used without limitation. SEB modifications may be prepared, for example, by producing SEB-producing cells with the genetic engineering technique and separating therefrom SEB modifications produced.

10 For example, wild type SEB (i.e. SEB that is prepared with the genetic engineering technique and has the same amino acid sequence as that of SEB from Staphylococcus aureus), from which SEB modifications for use in the medicament of the present invention can be deduced, is prepared as outlined below.

15 Since chromosomal DNA of SEB is known (Ranelli D. M. et al., Proc. Natl. Acad. Sci. USA, Vol. 82, p.5850- (1985)), 5'-sense and 3'-antisense primers can be prepared with a DNA synthesizer etc. Plaque hybridization is carried out with these primers and chromosomal DNAs present in the commercially available SEB DNA library to give plaques. PCR is then carried out with the sense primer and DNAs are extracted from the formed bands. The extracted DNAs are then inserted into an appropriate cloning vector for cloning.

25 Cloned SEB-coding gene is cleaved with restriction

enzymes such as SacI, HindIII, EcoRI, BamHI, XbaI, SalI and PstI and the obtained fragments are incorporated into a vector cleaved with restriction enzymes such as XmnI, HindIII, EcoRI, BamHI, XbaI, SalI and PstI to obtain recombinant DNA (Sambrook et al., Molecular Cloning, 2nd Ed., Chapter 9, 1989, New York, Cold Spring Harbor Laboratory Press). As a vector, expression vector for secretion pTrc99A etc. may suitably be employed.

The obtained recombinant DNA may be introduced into an appropriate host such as E. coli to give a transformant. After culturing said transformant in a usual manner, cells were separated after completion of culture. The separated cells were then ruptured in a usual manner and desired wild type SEB may be obtained from the suspension. Under suitable conditions, wild type SEB is conveniently secreted out into culture supernatant. In this case, the culture supernatant may be used as a starting material. A starting material is subjected to purification with a purification means such as, for example, immunoaffinity chromatography with absorbent to which anti-SEB monoclonal antibody is bound. A buffer used for final preparation in various experiments includes preferably Tris-HCl buffer, phosphate buffer, etc.

SEB modifications of the present invention may be prepared similarly as in preparation of wild type SEB as

mentioned above provided that binding sites with class II MHC molecule is analyzed based on data of X-ray structural analysis of SEB, specific sites of the amino acid sequence to be modified is determined, appropriate primers are prepared and point mutations are introduced via PCR.

For maintaining the activity of wild type SEB or SEB modifications prepared, freshly prepared one is used or if stored at 4°C, it is preferably used within about five days after storage. Alternatively, SEB modifications of the present invention may be stored under suitable circumstances of gelatin, salts, sugars, sugar alcohols or amino acids etc.

In accordance with the present invention, SEB modifications or derivatives thereof may be combined with known suitable excipients and formulated into prophylactics/remedies for immunopathy of the present invention in the known manner. A dosage form of the medicament of the present invention may be any as far as it is suited for oral administration, such as in the form of powder (solid), solution or syrup. For example, in a preferable embodiment, SEB modifications or derivatives thereof are lyophilized together with a suitable excipient, for example, carbohydrates, sugars, sugar alcohols, or amino acids etc. to produce a solid preparation. In another preferable embodiment, SEB modifications or derivatives thereof are dissolved in physiological saline or in a

suitable buffer with an ionic strength of acceptable tonicity to produce a liquid preparation. It is also conceivable that SEB modifications or derivatives thereof are dissolved in commercially available drink water and the solution is orally ingested. For a content in medicaments, SEB modifications or derivatives thereof are preferably present in an amount ranging from 0.05 μg to 50 mg (0.001 to 1,000 $\mu\text{g}/\text{kg}$ body weight), preferably from 0.5 μg to 0.5 mg (0.01 to 10 $\mu\text{g}/\text{kg}$ body weight) per administration.

An effective dose of prophylactics/remedies for immunopathy comprising as an active ingredient SEB modifications or derivatives thereof may vary depending on, for example, age of subject to be treated, symptoms or severity etc. and may finally be determined at physician's discretion. For example, the effective dose may typically be in a range of from 0.05 to 50,000 $\mu\text{g}/\text{day}$ for adult calculated as SEB modifications, and preferably 0.5 to 500 μg in total may be orally administered once or twice a day. SEB modifications of the present invention may optionally be combined with other medicaments such as, for example, azathioprine, cyclophosphamide, or high molecular weight antiinflammatory agents such as anti-TNF α antibody.

Major immunopathy that can be cured with prophylactics/medicaments for immunopathy comprising SEB modifications of the present invention is autoimmune disease

such as rheumatoid arthritis or ulcerative colitis. In addition to autoimmune disease, however, the prophylactics/medicaments for immunopathy of the present invention may also be applied to graft-versus-host disease after transplantation of bone marrow or type I, II, or III allergic diseases.

A "derivative" of SEB modifications of the present invention as used herein means the SEB modifications with substitution of at least one amino acid residues as mentioned above which are further modified. The present invention encompasses those SEB derivatives that have substitution, deletion or insertion at any amino acid residue within the amino acid sequence of natural type SEB in addition to the above specific amino acid residues in SEB modifications and still have an activity equivalent to that of SEB modifications.

INDUSTRIAL APPLICABILITY

In accordance with the present invention, novel prophylactics/remedies for immunopathy including autoimmune diseases such as rheumatoid arthritis, provision of which having been earnestly desired over a long period of term, can be provided.

EXAMPLE

The present invention is illustrated in more detail by means of the following Preparation and Examples,

but should not be construed to be limited thereto.

Preparation 1 (Preparation and expression of recombinant SEB modifications)

1-1 Cloning of SEB gene

5 A DNA library of Staphylococcus aureus enterotoxin A+B+D was purchased from CLONOTECH and plaque hybridization was carried out. As a probe, synthetic antisense DNA or PCR fragments were employed. A primer was added with SalI cleavage site at both ends for facilitating the subsequent
10 cloning procedure.

 Those plaques to which the primer bound were collected. PCR was carried out with a sense primer and DNA was extracted from the obtained bands and cloned into PCR-II vector (Invitrogen). The primers used in the above
15 hybridization are depicted in Table 1.

Table 1

Antisense: 5'-AAG TCG ACA ATA TTA GAA AAG GCA GGT ACT-3'

(SEQ ID NO: 1) SalI

Sense: 5'-ATG TCG ACT TAA TTG AAT ATT TAA GAT TAT-3'

20 (SEQ ID NO: 2) SalI

 Subsequently, a nucleotide sequence of the DNA was determined with an automatic sequencer. The obtained SEB gene contained a promoter region (SEB-Pro). In order to obtain SEB gene without the promoter region, PCR was further
25 conducted with primers listed in Table 2 and the obtained

DNA fragments were cloned into PCR-II vector.

Table 2

Sense: 5'-AAG TCG ACA AAA AAT GTA TAA GAG ATT ATT-3'

(SEQ ID NO: 3) SalI

5 Antisense: 5'-AAG TCG ACT TTC ACT TTT TCT TTG TCG TAA-3'

(SEQ ID NO: 4) SalI

A nucleotide sequence of the DNA of the obtained SEB gene was determined with an automatic sequencer. It was found that the thus obtained SEB gene did not contain any mutation. The SEB gene without the promoter region was cleaved with SalI and cloned into an expression vector for secretion pTrc99A (Pharmacia Biotech) with the same cleavage site. Using the vector where the gene was inserted at the right direction, expression was induced with IPTG to confirm expression and secretion of SEB.

1-2 Polymerase chain reaction (PCR)

PCR was carried out with Taq polymerase and DNA thermal cycler of Perkin Elmer Cetus (Norwalk, CT, U.S.A.) as described in Saiki et al. (Science vol. 239, p.487 (1988)). Each cycle consisted of a denaturing process (94°C for 1 minute) for denaturing and releasing a duplicate template DNA, an annealing process (55°C for 2 minutes) for annealing the primer and the template, and an elongation process (72°C for 2 minutes) for synthesis. A total of 30 to 35 cycles were repeated. The template was present at a

concentration of 1 to 10 M and the oligonucleotide primer at a concentration of 1 mM. A concentration of dNTP was 200 mM but was reduced to 20 mM for one dNTP when mutation is introduced.

5 1-3 Preparation and expression of recombinant SEB modification

Only SEB modifications with substitution of amino acid residues were prepared by the genetic recombination technique and expressed. SEB was modified for its region
10 involved in binding to MHC class II molecule as taught by Marrack P. et al. (J. Exp. Med., Vol., 171, p.445 (1990)). Table 3 indicates sites where substitution of amino acid residues was introduced.

Table 3

15	<u>Sites to be</u>	<u>Alteration</u>	<u>Indication</u>
	<u>modified</u>		
	D9	Asn → Asp	D9N
	F17	Phe → Ser	F17S
	N23	Asn → Asp	N23D
20	N23	Asn → Tyr	N23Y
	N23	Asn → Ile	N23I
	N23	Asn → Lys	N23K
	D9, N23	Asp → Asn	D9N N23D
		Asn → Asp	
25	F44	Phe → Ser	F44S

F44, I53	Phe → Leu	F44L I53N
	Ile → Asn	
F44, I53, G59	Phe → Leu	F44L I53N G59W
	Ile → Asn	
5	Gly → Trp	
I41, F44	Ile → Arg	I41R F44V
	Phe → Val	
I41, L45	Ile → Thr	I41T L45V
	Leu → Val	

10 1-4 Introduction of amino acid substitution

The SEB-Pro gene inserted into PCRII vector was used as a template DNA and SfiI and NotI sites were introduced at the 5'- and 3'-ends, respectively, by PCR. The obtained DNA fragment was inserted into pTrc99ApelB
 15 vector and a nucleotide sequence was determined by sequencing. Each one primer at the sense and antisense sites for PCR were prepared bearing mutation within the nucleotide sequence so that natural amino acid residues at the desired sites are altered to desired amino acid residues
 20 for each of SEB modifications. The primers at the antisense site (D9N, N23I, N23Y, N23D, N23K, F17S, F17S N23D)-SEB were firstly employed for preparation. Then, PCR was conducted using a primer for introducing SfiI site as a sense primer.

Figs. 1 and 2 show nucleotide sequences (SEQ ID
 25 NOS: 5 to 19) of PCR primers used for preparing SEB

modifications.

Subsequently, PCR was carried out using a primer for introducing NotI site, prepared previously using a sense primer, as an antisense primer. PCR was then carried out using the obtained two DNA fragments. The obtained DNA fragments were cloned into pTrc99A. The vector was then treated with SfiI and NotI to investigate whether or not the vector was cleaved into DNA fragments with desired size. Those DNA fragments with desired size were sequenced for their nucleotide sequence to confirm that mutation was correctly introduced.

1-5 Expression of SEB modifications and preparation of said modifications

SEB modifications were expressed with modification genes inserted into pTrc99A vector. E. coli cells with the gene incorporated were cultured in culture medium in which 4% CIRCLEGROW (BIO IOI Inc., Vista, CA, U.S.A.) and ampicillin (50 mg/ml) were dissolved at 37°C for 18 hours. The cells were collected and then suspended in the same medium to 0.3-1.0 of O.D. 550 nm. Thereto was added 2 mM isopropyl-B-D(-)-thiogalactopyranoside (IPTG) and expression was induced by shaking the mixture at 37°C overnight. After induction, the host E. coli cells were removed by centrifugation and supernatant was filtered through 0.45 µm filter membrane.

The thus prepared supernatant was passed through Sepharose 4B column to which anti-SEB monoclonal antibody SA58-2-6IgG was immobilized so that SEB modifications in the supernatant were adsorbed. The column was washed with 0.1 M Tris HCl (pH 8.0) and then eluted with 4M MgCl₂. The eluted fractions were dialyzed against 20-fold volume of physiological saline three times and then against 20-fold volume of PBS twice. All the SEB modifications prepared herein could be purified with the above monoclonal antibody column.

Example 1 (Lethal toxicity test with mice)

Natural type SEB does not provide mice with lethal toxicity. However, it is known that when D-galactosamine was previously administered, mice could be lead to death by intravenously or intraperitoneally administering 20 µg/mice of SEB as reported by Miethke T. et al. (J. Exp. Med. Vol., 175, p.91-98 (1992)). Thus, in order to investigate whether or not SEB modifications could reduce lethality, SEB or SEB modifications were administered to mice that previously received D-galactosamine.

Firstly, sensitivity to endotoxin was investigated. After administration of 20 mg/mice D-galactosamine to BALB/c mice, LPS (lipopolysaccharide) from E. coli B4 strain was intravenously administered to the mice and lethality after 24 hours was determined. As a result, death was not

observed with the dose of not more than 1 ng/mice of LPS (Table 4).

Table 4

	<u>Dose</u>	<u>Death No./Total No.</u>
5	1 µg/head	7/9
	100 ng/head	8/9
	10 g/head	5/9
	1 ng/head	0/9
	0.1 ng/head	0/9
10	An endotoxin level in samples of SEB modifications was reduced so that a final dose becomes not more than 10 ng/mice and male mice were used for an experiment. In case of male mice, lethal toxicity of SEB after administration of D-galactosamine was not less than	
15	100 µg/mice. Thus, SEB modifications were administered at a dose of 100 µg/mice. As shown in Table 5, natural type SEB, wild type SEB and D9N-SEB exhibited higher lethality, indicating lethal toxicity at that dose. In another SEB modifications, however, lethal toxicity was much reduced.	
20	"Natural type SEB" as used herein means enterotoxin from <u>Staphylococcus aureus</u> . "Wild type SEB" as used herein means SEB that has the same amino acid sequence as natural type SEB and is prepared by the genetic engineering technique.	

Table 5

<u>SEB modifications</u>		<u>Lethality (total death</u>	
<u>(100 µg/head)</u>		<u>No./ total No.) after:</u>	
		<u>24 hours</u>	<u>48 hours</u>
5	Natural type	8/10	8/10
	Wild type	7/10	9/10
	D9N	6/10	7/10
	I41T L45V	3/10	3/10
	I41R F44V	0/10	0/10
10	F44L I53N	0/9	0/9
	F44L I53N G59W	0/10	0/10
	F44S	0/10	0/10
	N23I	0/10	0/10
	N23D	0/10	0/10
15	N23Y	0/10	0/10
	N23K	0/10	0/10
	PBS	0/10	0/10

The same experiment was also carried out with female BALB/c mice. In case of female mice, high lethal toxicity was observed with 20 µg/mice SEB when 40 mg/mice D-galactosamine was administered. Higher lethality resulted in case of natural type SEB, wild type SEB and D9N-SEB. However, another SEB modifications exhibited lower lethality to demonstrate that lethal toxicity was reduced.

Example 2 (Comparison of TNF- α production by human peripheral blood mononuclear cells (PBMC) after stimulation with SEB or SEB modifications)

Peripheral blood mononuclear cells (PBMC) were collected from healthy adult using a lymphocyte separating reagent (Pharmacia) and suspended in RPMI1640 medium containing 10% FCS to make 1×10^6 cells/ml. The cell suspension (100 ml) was added to 96-well microtiter plate (Falcon) and then a solution of SEB or SEB modifications (100 ml) was added and the plate was incubated at 37°C for 24 hours. After 24 hours, culture supernatant was collected and TNF- α in the supernatant was quantitated with an enzyme immunoassay kit for human TNF- α (Biosource). The results are shown in Fig. 3. When PBMC was stimulated with SEB, a large amount of TNF- α was produced. On the contrary, when PBMC was stimulated with SEB modifications, TNF- α production was drastically reduced. Among these, N23Y showed the lowest, i.e. one-thousandth or less of specific activity as compared to SEB.

Example 3 (Inhibitory activity to T cells by SEB modifications)

Inhibitory activity to T cells by SEB modifications was estimated with purified peptide derivative (PPD)-responsive mouse T cell line PPD914. This T cell line is V β 8 TCR positive and can be responsive not

only to its antigen PPD but also to SEB. 1×10^4 PPD914 and 5×10^5 mitomycin-treated spleen cells from DBA/1J were cultured on 96-well microtiter plate (Falcon) and stimulated with addition of 0.5 mg/ml PPD. SEB modifications (1 mg/ml) were added to this system and culture was continued at 37°C for 48 hours. After 48 hours, tritium thymidine was added at 0.5 mCi/well and culture was further continued for 16 hours and proliferation was determined. Inhibitory activity of SEB modifications was estimated by comparing with the results obtained in the absence of SEB modifications. The results are shown in Fig. 4. SEB modifications N23Y and F44S inhibited T cell proliferation induced by its antigen PPD.

Example 4 (Estimation of SEB modifications for their safety in collagen-induced arthritis (CIA))

Collagen-induced arthritis (CIA) was induced in DBA/1 mice by intradermally administering 200 µg/mice of bovine type II collagen together with Freund's complete adjuvant to the mice. Twenty one days after administration of bovine type II collagen, each 50 µg/mice of SEB, F44S or N23D was intraperitoneally administered to the mice. A week later (28 days after the initial administration of collagen), onset rate of arthritis was compared. Table 6 summarizes the results of onset rate and severity of arthritis. Severe arthritis was observed in five among

eight mice administered with SEB. On the contrary, no arthritis could be found in mice administered with F44S or N23D.

Table 6

5	<u>Primary immunization</u>	<u>Booster</u>	<u>Onset rate</u>	<u>Severity</u>
	Collagen	SEB	5/8	6.2
	Collagen	F44S	0/8	-
	Collagen	N23D	0/8	-

Example 5 (Prophylactic effect to SEB-induced arthritis by
10 SEB modifications)

Each 50 µg/mice of F44S or N23D and phosphate
buffer as control was intraperitoneally administered to the
mice collagen-induced as in Example 4 eighteen days after
administration of bovine type II collagen. Three days
15 later, 50 µg/mice of SEB was intraperitoneally administered
to the mice. As shown in Table 7, arthritis was observed
in four among eight mice administered with phosphate buffer
(onset rate of 50%). On the contrary, onset rate was
reduced to 13% and 25% by previous administration of F44S
20 or N23D, respectively.

<u>Primary</u>	<u>Pretreatment</u>	<u>Booster</u>	<u>Onset_rate</u>	<u>Severity</u>
immunization				

Example 6 (Curative effect of SEB modifications via oral administration on collagen-induced arthritis (CIA))

20 After division into groups, an experiment started wherein mice were allowed to drink water unrestrictedly to thereby take SEB modifications and reduction of disease was observed. The disease of each mouse was scored once a week and a change in the severity was recorded. These results
25 are shown in Figs. 5 and 6. As to reduction of CIA

symptoms and inhibitory activity to exacerbation of CIA, I41R F44V, I41T L45V, D9N, N23K, F44S, N23I and wild type SEB were shown to be effective. Based on arthritis score in each experimental group, a test of significance was conducted with Wilcoxon test between the groups administered with PBS (the group with onset of CIA) and with SEB modifications. As a result, it was found that all the above SEB modifications were significant with a risk of less than 0.05 during 25 to 75 days after the onset of disease and hence could alleviate the symptoms of CIA. Thus, efficacy of SEB modifications that have lowered toxicity and retain efficacy to CIA was demonstrated, suggesting that these SEB modifications would be a potential medicament for human rheumatoid arthritis for oral administration with much lowered adverse side effects.

On the other hand, D9N N23D, F17S E22D and N23Y F208L were not significant with a risk of 0.05 in Wilcoxon test in comparison with the PBS-administered group.

Example 7 (Prophylactic effect of SEB modifications via oral administration on collagen-induced arthritis (CIA))

As described in Example 4, 200 µg of bovine type II collagen suspended in CFA was intradermally administered to DBA/I mice at the tail for immunization. Twenty days after the immunization, each 0.5 mg/mouse of SEB modifications dissolved in physiological saline was orally

10

WHAT IS CLAIMED IS:

1. A prophylactic/remedy for immunopathy comprising, as an active ingredient, modifications of Staphylococcal enterotoxin B (SEB) with substitution of at least one amino acid residues within the amino acid sequence of natural type SEB, or derivatives thereof, wherein said SEB modifications or derivatives thereof have inhibitory activity on T cell activation wherein they interact with specific V β component of T cell receptor (TCR) but are reduced in their immunological responsiveness to SEB without inducing elimination of T cells having specific V β component, the elimination being normally induced by natural type SEB or recombinant wild-type SEB.

2. The prophylactic/remedy for immunopathy of claim 1 wherein said SEB modifications or derivatives thereof are SEB having amino acid substitution at the 9-position on the basis of natural type SEB with a protein-constituting amino acid other than aspartic acid, or derivatives thereof.

3. The prophylactic/remedy for immunopathy of claim 2 wherein said SEB modifications or derivatives thereof are SEB having amino acid substitution at the 9-position on the basis of natural type SEB with asparagine.

4. The prophylactic/remedy for immunopathy of claim 1 wherein said SEB modifications or derivatives

thereof are SEB having amino acid substitution at the 23-position on the basis of natural type SEB with a protein-constituting amino acid other than asparagine, or derivatives thereof.

5 5. The prophylactic/remedy for immunopathy of claim 4 wherein said SEB modifications or derivatives thereof are SEB having amino acid substitution at the 23-position on the basis of natural type SEB with tyrosine, aspartic acid, isoleucine or lysine, or derivatives thereof.

10 6. The prophylactic/remedy for immunopathy of claim 1 wherein said SEB modifications or derivatives thereof are SEB having amino acid substitution at the 41-position on the basis of natural type SEB with a protein-constituting amino acid other than isoleucine plus amino
15 acid substitution either at the 44-position with a protein-constituting amino acid other than phenylalanine or at the 45-position with a protein-constituting amino acid other than leucine, or derivatives thereof.

20 7. The prophylactic/remedy for immunopathy of claim 6 wherein said SEB modifications or derivatives thereof are SEB having amino acid substitution at the 41-position on the basis of natural type SEB with arginine or threonine plus amino acid substitution either at the 44-position with valine or at the 45-position with valine, or
25 derivatives thereof.

8. The prophylactic/remedy for immunopathy of claim 1 wherein said SEB modifications or derivatives thereof are SEB having amino acid substitution at the 44-position on the basis of natural type SEB with a protein-constituting amino acid other than phenylalanine, or derivatives thereof.

9. The prophylactic/remedy for immunopathy of claim 8 wherein said SEB modifications or derivatives thereof are SEB having amino acid substitution at the 44-position on the basis of natural type SEB with serine, or derivatives thereof.

10. The prophylactic/remedy for immunopathy of claim 9 which is for oral administration.

11. The prophylactic/remedy for immunopathy of claim 10 which is an aqueous solution having pH and ionic strength that are physiologically acceptable.

12. The prophylactic/remedy for immunopathy of claim 11 wherein a substance selected from the group consisting of carbohydrates, sugars, sugar alcohols and amino acids is present in an amount sufficient for providing the solution with a physiologically acceptable tonicity.

13. The prophylactic/remedy for immunopathy of any one of claims 1 to 12 which is used for prophylactic/remedy of rheumatoid arthritis.

<110> JURIDICAL FOUNDATION THE CHEMO-SERO-THERAPEUTIC
RESEARCH INSTITUTE

5)

<150> JP 10-50137

<160> 19

10

<212> DNA

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20

<212> DNA

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by PCR amplification

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10 <400> 11

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<211> 42

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25 <220>

<223> Antisense primer designed for preparation of SEB variant by PCR amplification

<400> 17

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30 <210> 18

<211> 60

<220>

5

aaatctatag atcaactgct atactttgac ttaatatatt ctaacaagga cactaagtta 60

<211> 60

10

<223> Antisense primer designed for preparation of SEB variant by PCR amplification

taacttagtg tccttgttag aatatattaa gtcaaagtat agcagttgat ctatagattt 60

ABSTRACT OF THE DISCLOSURE

A prophylactic/remedy for immunopathy for immunopathy comprising, as an active ingredient, modifications of Staphylococcal enterotoxin B (SEB) with substitution of at least one amino acid residues within the amino acid sequence of natural type SEB, or derivatives thereof, wherein said SEB modifications or derivatives thereof have inhibitory activity on T cell activation wherein they interact with specific V β component of T cell receptor (TCR) but are reduced in their immunological responsiveness to SEB without inducing elimination of T cells having specific V β component, the elimination being normally induced by natural type SEB or recombinant wild-type SEB.

Fig. 1

D 9 N-S E B

Antisense: CTC GCG GCC CAG CCG GCC ATG GCC GAG AGT CAA CCA GAT CCT
 AAA CCA AAC GAG TTG CAC AAA TCG

N 23 I-S E B

Antisense: CTC GCG GCC CAG CCG GCC ATG GCC GAG AGT CAA CCA GAT CCT
 AAA CCA GAT GAG TTG CAC AAA TCG AGT AAA TTC ACT GGT TTG
 ATG GAA ATT ATG AAA GTT TT

N 23 K-S E B

Antisense: CTC GCG GCC CAG CCG GCC ATG GCC GAG AGT CAA CCA GAT CCT
 AAA CCA GAT GAG TTG CAC AAA TCG AGT AAA TTC ACT GGT TTG
 ATG GAA AAA ATG AAA GTT TT

N 23 Y-S E B

Antisense: CTC GCG GCC CAG CCG GCC ATG GCC GAG AGT CAA CCA GAT CCT
 AAA CCA GAT GAG TTG CAC AAA TCG AGT AAA TTC ACT GGT TTG
 ATG GAA TAT ATG AAA GTT TT

N 23 D-S E B

Antisense: CTC GCG GCC CAG CCG GCC ATG GCC GAG AGT CAA CCA GAT CCT
 AAA CCA GAT GAG TTG CAC AAA TCG AGT AAA TTC ACT GGT TTG
 ATG GAA GAT ATG AAA GTT TT

D 9 N N 23 D-S E B

Antisense: CTC GCG GCC CAG CCG GCC ATG GCC GAG AGT CAA CCA GAT CCT
 AAA CCA AAC GAG TTG CAC AAA TCG AGT AAA TTC ACT GGT TTG
 ATG GAA GAT ATG AAA GTT TT

Fig. 2

F 17 S-S E B

Antisense : CTC GCG GCC CAG CCG GCC ATG GCC GAG AGT CAA CCA GAT CCT
AAA CCA GAT GAG TTG CAC AAA TCG AGT AAA TCC ACT GGT TTG
ATG GAA

F 44 S-S E B

Sense : AAA TCT ATA GAT CAA TCT CTA TAC TTT GAC TTA
Antisense : TAA GTC AAA GTA TAG AGA TTG ATC TAT AGA TTT

I 41 T L 45 V-S E B

Sense : TCT ATA GAT CAA TTT GTA TAC TTT GAC TTA ATA
Antisense : TAT TAA GTC AAA GTA TAC AAA TTG ATC TAT AGA

I 41 R F 44 V-S E B

Sense : ATA AAC GTT AAA TCT CGT GAT CAA GTT CTA TAC TTT GAC TTA
Antisense : TAA GTC AAA GTA TAG AAC TTG ATC ACG AGA TTT AAC GTT TAT

F 44 L I 53 N-S E B

Sense : AAA TCT ATA GAT CAA CTG CTA TAC TTT GAC TTA ATA TAT TCT
AAC AAG GAC ACT AAG TTA
Antisense : TAA CTT AGT GTC CTT GTT AGA ATA TAT TAA GTC AAA GTA TAG
CAG TTG ATC TAT AGA TTT

Fig. 3

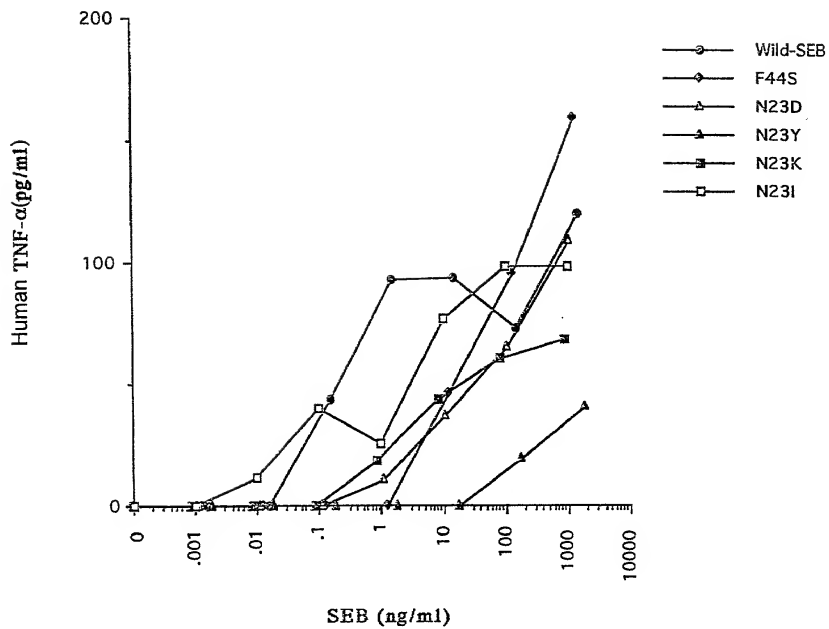


Fig. 4

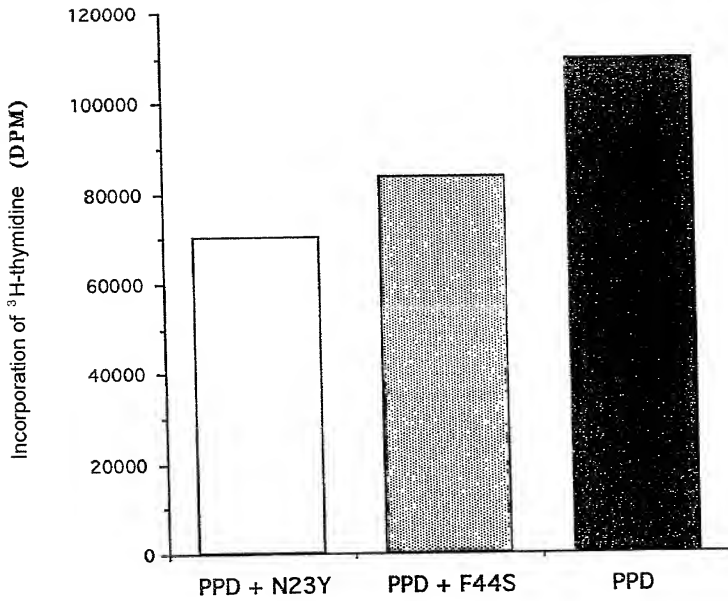


Fig. 5

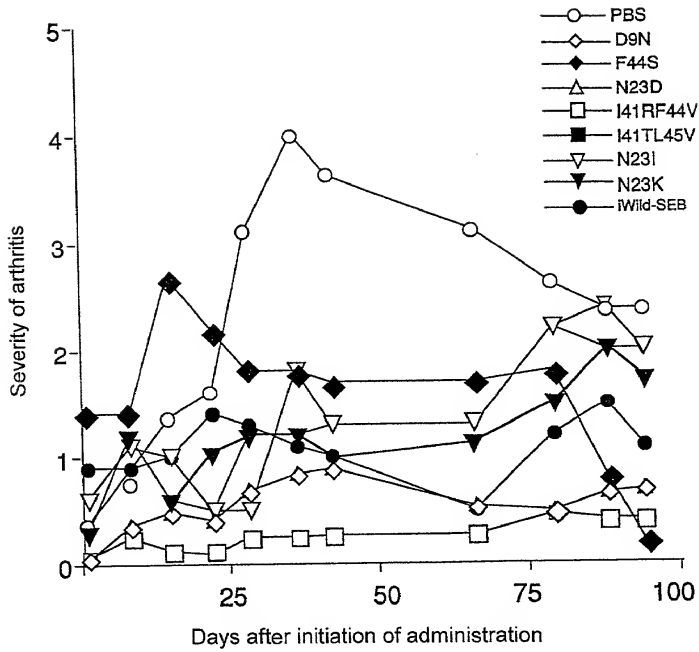
1. Effective cases

Fig. 6

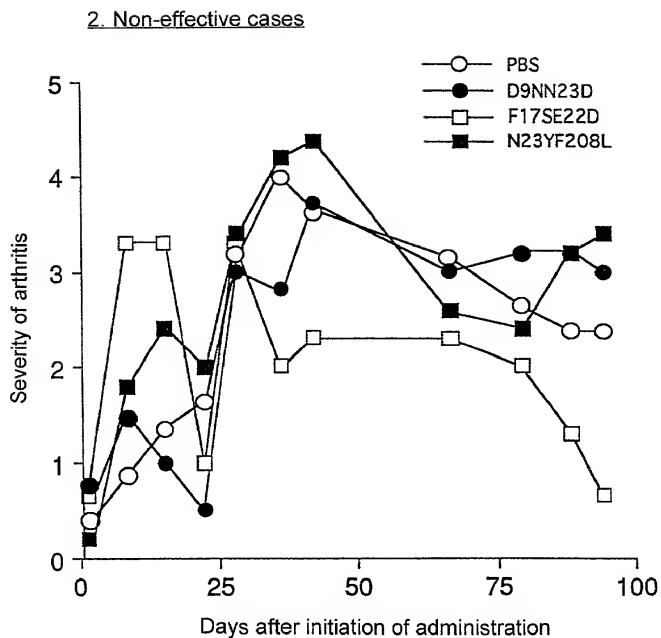
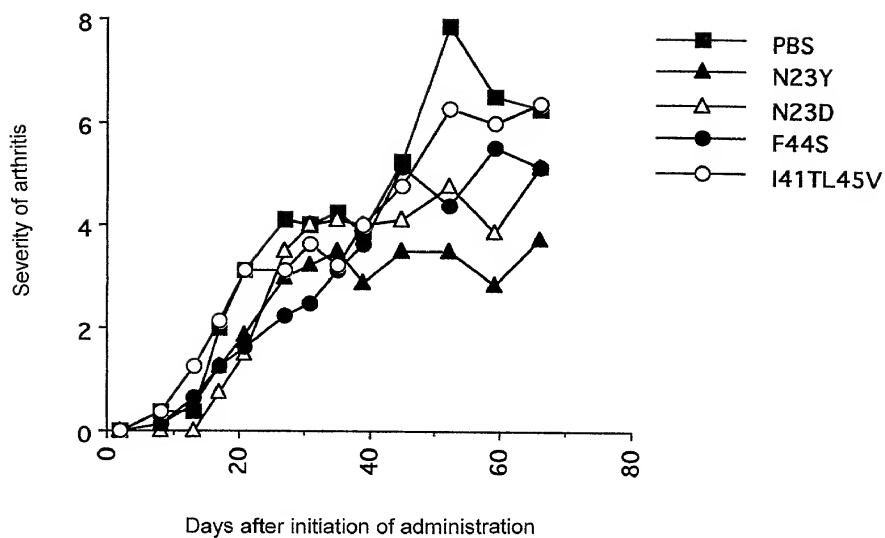


Fig. 7



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ATTORNEY DOCKET NO.
20-4736P

PLEASE NOTE:
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COMBINED DECLARATION AND POWER OF ATTORNEY FOR PATENT AND DESIGN APPLICATIONS

As a below named inventor, I hereby declare that: my residence, post office address and citizenship are as stated next to my name; that I verily believe that I am the original, first and sole inventor (if only one inventor is named below) or an original, first and joint inventor (if plural inventors are named below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

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the specification of which is attached hereto. If not attached hereto,

the specification was filed on _____ as
United States Application Number _____;
and amended on _____ (if applicable); and/or
the specification was filed on February 15, 1999 as PCT
International Application Number PCT/JP99/00638;
amended under PCT Article 19 on _____ (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I do not know and do not believe the same was ever known or used in the United States of America before my or our invention thereof, or patented or described in any printed publication in any country before my or our invention thereof or more than one year prior to this application, that the same was not in public use or on sale in the United States of America more than one year prior to this application, that the invention has not been patented or made the subject of an inventor's certificate issued before the date of this application in any country foreign to the United States of America on an application filed by me or my legal representatives or assigns more than twelve months (six months for designs) prior to this application, and that no application for patent or inventor's certificate on this invention has been filed in any country foreign to the United States of America prior to this application by me or my legal representatives or assigns, except as follows.

I hereby claim foreign priority benefits under Title 35, United States Code, §119 (a)-(d) of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)

Priority Claimed

<u>050137/1998</u> (Number)	<u>Japan</u> (Country)	<u>February 15, 1998</u> (Month / Day / Year Filed)	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> No <input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Month / Day / Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> No <input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Month / Day / Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> No <input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Month / Day / Year Filed)	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> No <input type="checkbox"/> No

Insert Priority
Information:
(if appropriate) →

Insert Provisional
Application(s): →
(if any)

I hereby claim the benefit under Title 35, United States Code, §119(e) of any United States provisional application(s) listed below.

_____ (Application Number)	_____ (Filing Date)
_____ (Application Number)	_____ (Filing Date)

All Foreign Applications, if any, for any Patent or Inventor's Certificate Filed More than 12 Months (6 Months for Designs) Prior to the Filing Date of This Application:

Insert Requested
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Country	Application Number	Date of Filing (Month / Day / Year)
_____	_____	_____
_____	_____	_____

I hereby claim the benefit under Title 35, United States Code, §120 of any United States and/or PCT application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States and/or PCT application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

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_____ (Application Number)	_____ (Filing Date)	_____ (Status — patented, pending, abandoned)
_____ (Application Number)	_____ (Filing Date)	_____ (Status — patented, pending, abandoned)

I hereby appoint the following attorneys to prosecute this application and/or an international application based on this application and to transact all business in the Patent and Trademark Office connected therewith and in connection with the resulting patent based on instructions received from the entity who first sent the application papers to the attorneys identified below, unless the inventor(s) or assignee provides said attorneys with a written notice to the contrary:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full Name of First or Sole Inventor:

Insert Name of Inventor →

Insert Date This Document is Signed →

Insert Residence →

Insert Citizenship →

Insert Post Office Address →

Address

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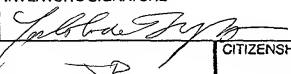
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see above

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Residence (City, State & Country)		CITIZENSHIP	
POST OFFICE ADDRESS (Complete Street Address including City, State & Country)			
GIVEN NAME	FAMILY NAME	INVENTOR'S SIGNATURE	DATE*
Residence (City, State & Country)		CITIZENSHIP	
POST OFFICE ADDRESS (Complete Street Address including City, State & Country)			

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see above

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see above

Full Name of Ninth
Inventor, if any:

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* DATE OF SIGNATURE